```
http://www.cas.org/legal/infopolicy.html
=> s (electrochemical cell) or battery
        166965 ELECTROCHEMICAL
            16 ELECTROCHEMICALS
        166980 ELECTROCHEMICAL
                 (ELECTROCHEMICAL OR ELECTROCHEMICALS)
        321556 ELECTROCHEM
            21 ELECTROCHEMS
        321563 ELECTROCHEM
                 (ELECTROCHEM OR ELECTROCHEMS)
        349261 ELECTROCHEMICAL
                 (ELECTROCHEMICAL OR ELECTROCHEM)
       2419216 CELL
       2087364 CELLS
       3164975 CELL
                 (CELL OR CELLS)
         13016 ELECTROCHEMICAL CELL
                 (ELECTROCHEMICAL (W) CELL)
        147421 BATTERY
        114715 BATTERIES
        160332 BATTERY
                 (BATTERY OR BATTERIES)
L1
        170768 (ELECTROCHEMICAL CELL) OR BATTERY
=> s l1 and (lithium (3a) air) or (li (3a) air)
        343773 LITHIUM
           372 LITHIUMS
        343901 LITHIUM
                 (LITHIUM OR LITHIUMS)
       1060546 AIR
           455 AIRS
       1060734 AIR
                 (AIR OR AIRS)
           599 LITHIUM (3A) AIR
        208015 LI
          1071 LIS
        208917 LI
                 (LI OR LIS)
       1060546 AIR
           455 AIRS
       1060734 AIR
                 (AIR OR AIRS)
           274 LI (3A) AIR
L2
           381 L1 AND (LITHIUM (3A) AIR) OR (LI (3A) AIR)
=> d his
     (FILE 'HOME' ENTERED AT 15:46:35 ON 08 JUL 2008)
     FILE 'REGISTRY' ENTERED AT 15:46:49 ON 08 JUL 2008
     FILE 'CAPLUS' ENTERED AT 15:46:53 ON 08 JUL 2008
L1
         170768 S (ELECTROCHEMICAL CELL) OR BATTERY
L2
            381 S L1 AND (LITHIUM (3A) AIR) OR (LI (3A) AIR)
=> s 12 and intercalat?
         50984 INTERCALAT?
            13 L2 AND INTERCALAT?
L3
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SOURCE:

(FILE 'HOME' ENTERED AT 15:46:35 ON 08 JUL 2008)

FILE 'REGISTRY' ENTERED AT 15:46:49 ON 08 JUL 2008

FILE 'CAPLUS' ENTERED AT 15:46:53 ON 08 JUL 2008

L1 170768 S (ELECTROCHEMICAL CELL) OR BATTERY

L2 381 S L1 AND (LITHIUM (3A) AIR) OR (LI (3A) AIR)

L3 13 S L2 AND INTERCALAT?

=> d 1-13 ibib ti ti abs

L3 ANSWER 1 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2008:342025 CAPLUS <<LOGINID::20080708>>

TITLE: Effect of catalyst on the performance of rechargeable

lithium/air batteries

AUTHOR(S): Debart, A.; Bao, J.; Armstrong, G.; Bruce, P. G. CORPORATE SOURCE: EaStCHEM, School of Chemistry, University of St

Andrews, North Haugh, St Andrews, Fife, KY16 9ST, UK ECS Transactions (2007), 3(27, Lithium-Ion Batteries),

 $\frac{1}{2}$

225-232

CODEN: ECSTF8

PUBLISHER: Electrochemical Society

DOCUMENT TYPE: Journal; (computer optical disk)

LANGUAGE: English

TI Effect of catalyst on the performance of rechargeable lithium/ air batteries

TI Effect of catalyst on the performance of rechargeable lithium/ air batteries

AB Rechargeable lithium batteries are now a major technol., driven by their superior energy d. compared with alternative rechargeable batteries. There is much interest in increasing further the energy d. This is limited by the pos. electrode, LiCoO2, which can cycle only around 0.5 Li per formula unit. Intensive research world-wide on new intercalation cathodes will increase the amount of Li that may be stored, but only by a factor of 2. Replacing the intercalation cathode with an O2 electrode could raise the capacity 5-10-fold. Here we explore the influence of the catalyst could have on the performance of a non-aqueous O2 electrode on a Li/O2 cell. Of the catalysts studied, the highest capacity retentions were observed with Fe3O4, CuO and CoFe2O4 (6.67% per cycle). Co3O4 gives the best compromise between initial capacity (2000mAhg-1) and capacity retention (6.5% per cycle), as well as the lowest charging voltage 4V.

REFERENCE COUNT: 23 THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 2 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2007:882099 CAPLUS <<LOGINID::20080708>>

TITLE: Rechargeable lithium/air

battery

AUTHOR(S): Debart, Aurelie; Armstrong, Graham; Bao, Jianli;

Paterson, Allan J.; Bruce, Peter G.

CORPORATE SOURCE: School of Chemistry, University of St Andrews, St

Andrews, KY169ST, UK

SOURCE: Abstracts of Papers, 234th ACS National Meeting,

Boston, MA, United States, August 19-23, 2007 (2007), FUEL-227. American Chemical Society: Washington, D.

С.

CODEN: 69JNR2

DOCUMENT TYPE: Conference; Meeting Abstract; (computer optical disk)

LANGUAGE: English

AB

SOURCE:

ΤT Rechargeable lithium/air battery

ΤТ Rechargeable lithium/air battery

Rechargeable lithium batteries already dominate consumer electronics and have a crucial role to play addressing global warming and powering medical devices. Their energy storage (d.) is limited by the quantity of Li that can be stored in the pos. intercalation electrode LiCoO2. To take a leap forward in energy storage, a radically

different approach is required e.g. replacing LiCoO2 by reaction with O2 from the air. Such an electrode can be repeatedly charged and discharged and the Li2O2 formed on discharge is electrochem. decomposed to Li + O2 on charge. Here we explore the influence of the catalyst could have on the

performance of a non-aqueous O2 electrode on a Li/O2 cell.

ANSWER 3 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER:

TITLE: Rechargeable lithium/air

battery

Debart, Aurelie; Armstrong, Graham; Bao, Jianli; AUTHOR(S):

Paterson, Allan J.; Bruce, Peter G.

CORPORATE SOURCE: EaStCHEM, School of Chemistry, University of St

> Andrews, North Haugh, St Andrews, Fife, KY16 9ST, UK Preprints of Symposia - American Chemical Society,

Division of Fuel Chemistry (2007), 52(2), 692

CODEN: PSADFZ; ISSN: 1521-4648

PUBLISHER: American Chemical Society, Division of Fuel Chemistry

DOCUMENT TYPE: Journal; (computer optical disk)

LANGUAGE: English ΤI Rechargeable lithium/air battery

ΤI Rechargeable lithium/air battery

AΒ Rechargeable lithium batteries already dominate consumer

electronics and have a crucial role to play addressing global warming and powering medical devices. Their energy storage (d.) is limited by the quantity of Li that can be stored in the pos. intercalation electrode LiCoO2. To take a leap forward in energy storage, a radically different approach is required e.g. replacing LiCoO2 by reaction with O2 from the air. Such an electrode can be repeatedly charged and discharged and the Li2O2 formed on discharge is electrochem. decomposed to Li + O2 on

charge. Here we explore the influence of the catalyst could have on the performance of a non-aqueous 02 electrode on a Li/O2 cell.

REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 4 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2004:312161 CAPLUS <<LOGINID::20080708>>

DOCUMENT NUMBER: 140:342143

TITLE: Secondary nonaqueous electrolyte air battery INVENTOR(S): Kuboki, Takashi; Okuyama, Akio; Osaki, Takahisa;

Takami, Norio

Toshiba Corp., Japan PATENT ASSIGNEE(S):

Jpn. Kokai Tokkyo Koho, 23 pp. SOURCE:

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2004119278 JP 4015916	A B2	20040415 20071128	JP 2002-283527	20020927

PRIORITY APPLN. INFO.: JP 2002-283527 20020927

TI Secondary nonaqueous electrolyte air battery TI Secondary nonaqueous electrolyte air battery

GΙ

AB The battery has a nonaq. electrolyte solution between a Liintercalating anode and a cathode in a battery case having air
inlet holes supplying O to the cathode; where the electrolyte solution
comprises an ordinary temperature molten salt, containing Li+ and ≥1 cation
selected from I (R1-4 = C<8 alkyl, C<8 Ph, or C<8 benzyl group) II (R5,
R7= C<8 alkyl, C<8 Ph, or C<8 benzyl group; and R6, R8= H, C<8 alkyl, C<8
Ph, or C<8 benzyl group) , and III (R9 = C<8 alkyl, C<8 Ph, or C<8 benzyl
group)

L3 ANSWER 5 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2004:65690 CAPLUS <<LOGINID::20080708>>

DOCUMENT NUMBER: 140:202347

TITLE: A better understanding of the irreversible lithium

insertion mechanisms in disordered carbons

AUTHOR(S): Beguin, F.; Chevallier, F.; Vix, C.; Saadallah, S.;

Rouzaud, J. N.; Frackowiak, E.

CORPORATE SOURCE: Centre de Recherche sur la Matiere Divisee,

CNRS-University, Orleans, 45071, Fr.

SOURCE: Journal of Physics and Chemistry of Solids (2004),

65(2-3), 211-217

CODEN: JPCSAW; ISSN: 0022-3697

PUBLISHER: Elsevier Science B.V.

DOCUMENT TYPE: Journal LANGUAGE: English

TI A better understanding of the irreversible lithium insertion mechanisms in disordered carbons

TI A better understanding of the irreversible lithium insertion mechanisms in disordered carbons

AB Disordered carbons have high Li storage capacities. However, compared to graphite, they generally show high hysteresis and irreversibility, and sometimes a poor cycleability, which strongly limit their practical application. To better understand the phenomena occurring during the 1st reduction (discharge) of nanostructured carbons and to improve electrochem. performance, the insertion reaction was studied. Irreversible capacity is related to the sp. surface area, but many deviations from this relationship are observed A better linear relation is obtained by considering the active surface area. Deactivating the surface by a pyrolytic C deposit allows the irreversible capacity to be reduced. The electrochem. properties of this new C/C composite were studied by galvanostatic cycling.

REFERENCE COUNT: 20 THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ACCESSION NUMBER: 2002:953986 CAPLUS <<LOGINID::20080708>>

DOCUMENT NUMBER: 138:306709

TITLE: CeO2 thin films obtained by sol-gel deposition and

annealed in air or argon

AUTHOR(S): Kozjek Skofic, Irena; Sturm, Saso; Ceh, Miran;

Bukovec, Natasa

CORPORATE SOURCE: Faculty of Chemistry and Chemical Technology,

University of Ljubljana, Ljubljana, SI-1000, Slovenia

SOURCE: Thin Solid Films (2002), 422(1-2), 170-175

CODEN: THSFAP; ISSN: 0040-6090

PUBLISHER: Elsevier Science B.V.

DOCUMENT TYPE: Journal LANGUAGE: English

TI CeO2 thin films obtained by sol-gel deposition and annealed in air or argon

TI CeO2 thin films obtained by sol-gel deposition and annealed in air or argon

AB Thin films of CeO2 were prepared on SnO2/F-coated glass plates by the sol-gel dip-coating process using CeCl3·7H2O as a precursor. The films were heat-treated in an air or argon atmospheric The structural, electrochem. and optical properties of these films depend on the preparation conditions. Transmission electron microscopy (TEM) showed the films to be polycryst. with randomly oriented crystallized domains of up to 10 nm in size. The degree of crystallinity of films heat-treated in argon is higher than that of those heat-treated in air, and therefore their charge capacity values (15.9 mC cm-2 after 100 cycles) and reversibility of the ion-storage process (0.99 after 100 cycles) are higher than for films heat-treated in air (10.5 mC cm-2 and 0.86 after 100 cycles, resp.). Both films are optically passive under Li+ ion insertion and have high optical transmittance (>80%).

REFERENCE COUNT: 22 THERE ARE 22 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 7 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2002:193273 CAPLUS <<LOGINID::20080708>>

DOCUMENT NUMBER: 136:234670

TITLE: Cathode active materials, their manufacture, and

secondary organic electrolyte batteries

using them

INVENTOR(S): Oikawa, Takao; Iwatani, Keizo; Yamaguchi, Masashi

PATENT ASSIGNEE(S): Chisso Corp., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 6 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2002075355	A	20020315	JP 2000-251956	20000823
PRIORITY APPLN. INFO.:			JP 2000-251956	20000823

- TI Cathode active materials, their manufacture, and secondary organic electrolyte batteries using them
- TI Cathode active materials, their manufacture, and secondary organic electrolyte batteries using them
- AB The cathode active materials (Li1-wZnw)8a(Mn2-x-yM1yLix)16d04 (M1 = A1, Co, and/or Ni; $0 < w \le 0.1$; x = 0-0.1; $0 < y \le 0.1$) for secondary batteries using Li-intercalatable carbonaceous materials for anodes and mixts. of organic solvents and Li salt electrolytes as electrolyte solns., are manufactured by mixing Al compds., Co

compds., and/or Ni compds., Mn2O3 prepared by heating Mn oxides at $550-750^{\circ}$, Zn compds., and Li compds., pressurizing the mixts. with air at O partial pressure 0.04-1 MPa, heating them at $600-900^{\circ}$ for 1-40 h, and gradually cooling them at a cooling rate of $\leq 10^{\circ}/\text{min to 500°}$. Secondary organic electrolyte batteries using the cathode active materials show long cycle life at high temperature

ANSWER 8 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER:

DOCUMENT NUMBER: 131:274097

TITLE: Effects of CO2 in air on Li

deintercalation from LiNi1-x-yCoxAlyO2

Matsumoto, Kazutoshi; Kuzuo, Ryuichi; Takeya, Kaname; AUTHOR(S):

Yamanaka, Atsushi

Central Research Laboratories, Sumitomo Metal Mining, CORPORATE SOURCE:

Ichikawa-shi, Chiba, Japan

Journal of Power Sources (1999), 81-82, 558-561 SOURCE:

CODEN: JPSODZ; ISSN: 0378-7753

Elsevier Science S.A. PUBLISHER:

DOCUMENT TYPE: Journal LANGUAGE: English

Effects of CO2 in air on Li deintercalation from

LiNi1-x-yCoxAlyO2

Effects of CO2 in air on Li deintercalation from TΙ

LiNi1-x-yCoxAlyO2

and

AUTHOR(S):

The reaction of CO2 and practical high-performance cathode materials AB LiNi1-x-yCoxAlyO2 was examined with XRD at RT and high temperature, TGA, IR,

chemical anal. Even at room temperature, Li deintercalation took place and formed

Li2CO3 on the mother surface. The conversion to Li2CO3 in air at 55% RH at 25° was in proportion to the square root of exposure time. It was 8% after being left for 500 h. Using high-temperature XRD at 25-800°, the Li2CO3 phase formation was confirmed from about 500° , and the conversion exceeded 70% under atmospheric CO2 at 675°. The activation energy of the reaction of deintercalated Li and CO2 was estimated by Ozawa's method. The activation energy obtained using Ozawa's method was 130 kJ/mol for CO2 diffusing through the Li2CO3 layer

which formed on the surface during the reaction. REFERENCE COUNT: THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 9 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN

1999:399403 CAPLUS <<LOGINID::20080708>> ACCESSION NUMBER:

DOCUMENT NUMBER: 131:90157

TITLE: Synthesis of LixNi0.85Co0.1502 by the PVA-precursor

> method and the effect of air flow during the pyrolysis Kweon, Ho-Jin; Kim, Geun Bae; Kim, Sue Joo; Song, Me Young; Park, Seon Hui; Kwon, Hye Young; Park, Dong Gon

CORPORATE SOURCE: Department of Chemistry, Sookmyung Women's University,

Seoul, 140-742, S. Korea

SOURCE: Bulletin of the Korean Chemical Society (1999), 20(5),

508-516

CODEN: BKCSDE; ISSN: 0253-2964

PUBLISHER: Korean Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English

Synthesis of LixNi0.85Co0.1502 by the PVA-precursor method and the effect ΤТ of air flow during the pyrolysis

ΤТ Synthesis of LixNi0.85Co0.1502 by the PVA-precursor method and the effect of air flow during the pyrolysis

Polycryst. powder of LixNi0.85Co0.1502 was synthesized by pyrolyzing a AΒ powder precursor obtained by the PVA-precursor method. Coin cells of lithium-ion rechargeable battery were assembled, whose cathodes were fabricated from the crystalline powders of LixNi0.85Co0.1502 synthesized by the method. The effect of synthetic variation on the property of the cell was tested by carrying out 100 consecutive cycles of charge-discharge on the cells. The property of the cell was largely influenced by the pyrolysis conditions applied for the synthesis of the LixNi0.85Co0.1502. Depending on whether the pyrolysis was carried out in standing air or in the flow of dry air, the discharge capacity and cycle-reversibility of the cell varied in large extent. When the powder precursor was pyrolyzed in standing air, a minor phase of lithium carbonate was remained in the LixNi0.85Co0.1502. The carbon containing powder precursor had to be pyrolyzed in the flow of dry air to eliminate the minor phase. In the flow of dry air, the lithium carbonate in the precursor was eliminated over 500-700° without any prominent heat event. By controlling the flow of air over the precursor during its pyrolysis, particle size could also be altered. The effect of flowing dry air, during first step pyrolysis or during second step heat treatment, on the property of the cell was discussed.

REFERENCE COUNT: 30 THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 10 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1998:198116 CAPLUS <<LOGINID::20080708>>

DOCUMENT NUMBER: 128:232838

ORIGINAL REFERENCE NO.: 128:46049a,46052a
TITLE: Air-lithium secondary

batteries with carbonaceous anodes

INVENTOR(S): Takami, Norio; Osaki, Takahisa

PATENT ASSIGNEE(S): Toshiba Corp., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 5 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PAT	TENT NO.	KIND	DATE	AP.	PLICATION NO.		DATE
						-	
JP	10083836	A	19980331	JΡ	1996-239156		19960910
JP	3699786	В2	20050928				
JP	2005166685	A	20050623	JΡ	2005-30487		20050207
PRIORITY	APPLN. INFO.:			JΡ	1996-239156	АЗ	19960910

TI Air-lithium secondary batteries with carbonaceous anodes

TI Air-lithium secondary batteries with carbonaceous anodes

AB The title batteries use anodes containing Li-intercalating carbonaceous materials and solid electrolytes. The batteries have high capacity and long cycle life.

L3 ANSWER 11 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1998:155031 CAPLUS <<LOGINID::20080708>>

DOCUMENT NUMBER: 128:219477

ORIGINAL REFERENCE NO.: 128:43441a,43444a

TITLE: Nonaqueous electrolyte secondary batteries

with high capacity and long cycle life and their

manufacture

INVENTOR(S): Tanaka, Mitsutoshi; Ishizuka, Hiroshi; Matsufuji,

Akihiro

PATENT ASSIGNEE(S): Fuji Photo Film Co., Ltd., Japan; Ube Industries, Ltd.

SOURCE: Jpn. Kokai Tokkyo Koho, 13 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 10064540	A	19980306	JP 1996-217531	19960819
JP 3663763	В2	20050622		

PRIORITY APPLN. INFO.: JP 1996-217531 19960819

TI Nonaqueous electrolyte secondary batteries with high capacity and long cycle life and their manufacture

TI Nonaqueous electrolyte secondary batteries with high capacity and long cycle life and their manufacture

AB The batteries have cathodes containing Li intercalatable compds., anodes containing amorphous chalcogen compds. and/or amorphous oxides as main components and ≥3 kinds of Group IA-IVA elements, Li salt-containing nonaq. electrolytes treated by air-bubbling, and separators. The nonaq. electrolytes are air-bubbled before injected into battery cases.

L3 ANSWER 12 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1998:86584 CAPLUS <<LOGINID::20080708>>

DOCUMENT NUMBER: 128:175259

ORIGINAL REFERENCE NO.: 128:34401a,34404a

TITLE: Structural in-situ study of Li intercalation

in Li1+ α Mn2- α O4 spinel-type oxides

AUTHOR(S): Le Cras, F.; Anne, M.; Bloch, Didier; Strobel, P.

CORPORATE SOURCE: Laboratoire de Cristallographie CNRS, BP 166, Grenoble

Cedex 9, 38042, Fr.

SOURCE: Solid State Ionics (1998), 106(1,2), 1-10

CODEN: SSIOD3; ISSN: 0167-2738

PUBLISHER: Elsevier Science B.V.

DOCUMENT TYPE: Journal LANGUAGE: English

TI Structural in-situ study of Li intercalation in

 $Li1+\alpha Mn2-\alpha O4$ spinel-type oxides

TI Structural in-situ study of Li intercalation in

 $Li1+\alpha Mn2-\alpha O4$ spinel-type oxides

AB Spinel oxides LiMn204 and Li1.23Mn1.7704 were studied using an air —tight electrochem. Li cell mounted on an x-ray powder diffractometer. The combination of electrochem. and x-ray data unambiguously shows the appearance of a tetragonal phase as soon as Li intercalation in the host oxide begins. The Li intercalation reaction is two-phase for both oxide hosts, in spite of the high initial Mn oxidation state in Li1.23Mn1.7704 (+3.82), which lies well above the expected Jahn-Teller distortion limit (+3.5). The tetragonal distortion is much stronger in lithiated Li2Mn204 than in Li1.23+xMn1.7704.

REFERENCE COUNT: 13 THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 13 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1991:683569 CAPLUS <<LOGINID::20080708>>

DOCUMENT NUMBER: 115:283569

ORIGINAL REFERENCE NO.: 115:48133a,48136a

TITLE: Lithium metal-free rechargeable batteries based on

lithium manganese oxide (Li1+xMn2O4) cathodes (0

 \leq + \leq 1) and carbon anodes Tarascon, J. M.; Guyomard, D.

CORPORATE SOURCE: Bellcore, Red Bank, NJ, 07701, USA

SOURCE: Journal of the Electrochemical Society (1991),

138(10), 2864-8

CODEN: JESOAN; ISSN: 0013-4651

DOCUMENT TYPE: Journal LANGUAGE: English

AUTHOR(S):

TI Lithium metal-free rechargeable batteries based on lithium manganese oxide (Li1+xMn2O4) cathodes (0 \leq + \leq 1) and carbon anodes

TI Lithium metal-free rechargeable batteries based on lithium manganese oxide (Li1+xMn2O4) cathodes (0 \leq + \leq 1) and carbon anodes

AB The spinel LiMn204 can be used as the cathode in rechargeable rocking-chair batteries based on Li intercalation anodes (carbon, either graphite or petroleum coke). At room temperature, such cells show promising cycle life, an average open-circuit voltage of 3.7 V and a specific energy of 250 W-h/kg of electrode materials (cathode + anode). A novel synthesis Li2Mn2O4, at <100° was developed, using LiI as a mild reducing agent. The cycling behavior of rocking-chair cells using the lithiated phase as the starting cathode is presented. Li2Mn2O4 appears to be a promising practical air stable Li -bearing cathode for rocking-chair-type rechargeable cells.

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